SEPARATION OF NUCLEOTIDES AND DNA FRAGMENTS BY CAPILLARY ELECTROPHORESIS USING WALL COATED CAPILLARIES.

<u>JOSHUA DEDERICH</u>, STEVEN A. STEINER, CHOUA YANG, JAMES P. HAMILTON, BRENDEN J. CARROLL, Department of Chemistry and Engineering Physics, University of Wisconsin-Platteville, Platteville WI 53818.

JAMES S. FRITZ, Department of Chemistry and Ames Laboratory, Iowa State University, Ames, IA 50011.

Capillary electrophoresis (CE) can be used to separate nucleotides and DNA fragments by using a capillary coated with Poly(diallyldimethyl-ammonium chloride) (PDADMAC). By coating the inside wall of the capillary with this cationic polymer a thin surface coating establishes a stable anodic electroosmotic flow. A simple capillary coating procedure was developed which consists of treating a new capillary by rinsing for 2 minutes with each of the following 0.1 M NaOH, purified water, 0.5% solution of PDADMAC and the background electrolyte (BGE) solution. This series of capillary rinses is adequate to provide a PDADMAC coating that has a wide pH range and can be used with simple BGE. A simpler BGE gives the potential for easier interfacing of CE instruments to more sensitive detectors like mass spectrometers and it also makes for a simpler analyte recovery after the separation. A systematic study of experimental parameters was done on synthetic mixtures of nucleic acids and DNA fragments to determine the optimum conditions for electrolyte concentration, pH and a suitable buffer for the BGE. Excellent separation of a mixture of four nucleotides in under 6 minutes were achieved. Electropherograms showing the separation attained for six DNA fragments from a sample of lambda DNA Ecor I Markers using PDADMAC coated columns will be shown.